

(FILE 'HOME' ENTERED AT 05:35:24 ON 21 MAR 2005)

FILE 'USPATFULL' ENTERED AT 05:35:29 ON 21 MAR 2005
ACTIVATE L10768359/L

L1 (10638)SEA FILE=USPATFULL ABB=ON PLU=ON TYROSINE AND (LIPOIC OR GLUT
L2 (360)SEA FILE=USPATFULL ABB=ON PLU=ON TYROSINE (100A) (LIPOIC OR G
L3 (174)SEA FILE=USPATFULL ABB=ON PLU=ON L2 AND (TOPICAL OR EXTERNAL)
L4 (198)SEA FILE=USPATFULL ABB=ON PLU=ON TYROSINE (50A) (LIPOIC OR GL
L5 (97)SEA FILE=USPATFULL ABB=ON PLU=ON L4 AND L3
L6 (89)SEA FILE=USPATFULL ABB=ON PLU=ON L5 NOT PERRICONE
L7 (97)SEA FILE=USPATFULL ABB=ON PLU=ON L5 NOT PERRICONE/AU
L8 (10)SEA FILE=USPATFULL ABB=ON PLU=ON (TYROSINE/CLM (50A) (LIPOIC/
L9 (83)SEA FILE=USPATFULL ABB=ON PLU=ON L6 NOT L8

L10 97 S L5
E PERIICONE CICHOMAS/AU
E PERIICONE NICHOLAS/AU
E PERRICONE NICHOLAS/AU
L11 45 S E27-29
L12 90 S L10 NOT L11
L13 90 FOCUS L12 1-
L14 4255 S DIMETHYLAMINOETHANOL
L15 174 S L3
L16 32 S DIMETHYLAMINOETHANOL/AB
L17 269 S DIMETHYLAMINOETHANOL/CLM
L18 273 S L16 OR L17
L19 8 S L18 AND L15
L20 4255 S DIMETHYLAMINOETHANOL
L21 50125 S TYROSINE
L22 26269 S (LIPOIC ACID OR GLUTATHIONE)
L23 23 S L20 (30A) L21 (30A) L22
L24 17 S L23 NOT L11
L25 8 S L20/CLM AND L21/CLM AND L22/CLM
L26 1 S L25 NOT L11
L27 7485 S L20/CLM OR L21/CLM OR L22/CLM
L28 2220 S L20/AB OR L21/AB OR L22/AB
L29 6 S L28 AND L27 AND L23
L30 0 S L29 NOT L11
L31 9 S L28 AND L27 AND L15
L32 2 S L31 NOT L11

=> save all
ENTER NAME OR (END) :end

=> save all temp
ENTER NAME OR (END) :l10768359/l
'L10768359/L' IN USE
A single name cannot be used for two saved items at the same time.
Enter "Y" if you wish to replace the current saved name with a new
definition. Enter "N" if the current saved definition must be
preserved. You may then reenter the SAVE command with a different
saved name. Enter "DISPLAY SAVED" at an arrow prompt (=>) to see a
list of your currently defined saved names.
REPLACE OLD DEFINITION? Y/(N):y
L# LIST L1-L32 HAS BEEN SAVED AS 'L10768359/L'

=>
Connection closed by remote host

EPRESENTATIVE: AUDLEY A. CIAMPORCERO JR., JOHNSON & JOHNSON, ONE
JOHNSON & JOHNSON PLAZA, NEW BRUNSWICK, NJ, 08933-7003
NUMBER OF CLAIMS: 24
EXEMPLARY CLAIM: 1
NUMBER OF DRAWINGS: 1 Drawing Page(s)
LINE COUNT: 822

L24 ANSWER 13 OF 17 USPATFULL on STN

SUMM . . . traps, retinoids such as retinol and retinyl palmitate, ceramides, polyunsaturated fatty acids, essential fatty acids, enzymes, enzyme inhibitors, minerals, estrogens, 2-dimethylaminoethanol, copper peptides such as Cu:GHK, lipoic acid, amino acids such a proline and tyrosine, lactobionic acid, acetyl-coenzyme A, niacin, riboflavin, thiamin, ribose, electron transporters such as NADH and FADH2; and botanical extracts such as.

ACCESSION NUMBER: 2002:279707 USPATFULL
TITLE: Composition containing Hedychium extract and use thereof
INVENTOR(S): Martin, Katharine M., Ringoes, NJ, UNITED STATES
Saliou, Claude, Gladstone, NJ, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002155138	A1	20021024
APPLICATION INFO.:	US 2002-52315	A1	20020118 (10)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2001-262822P	20010119 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	AUDLEY A. CIAMPORCERO JR., JOHNSON & JOHNSON, ONE JOHNSON & JOHNSON PLAZA, NEW BRUNSWICK, NJ, 08933-7003	
NUMBER OF CLAIMS:	20	
EXEMPLARY CLAIM:	1	
LINE COUNT:	517	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L24 ANSWER 14 OF 17 USPATFULL on STN

DETD . . . tocopheryl acetate; retinoids such retinol, retinal, retinyl palmitate, retinyl acetate, and retinoic acid; hormones such as estrogens and dihydroxyandrostene dione; 2-dimethylaminoethanol; lipoic acid; amino acids such a proline and tyrosine; lactobionic acid; self-tanning agents such as dihydroxy acetone; dimethyl aminoethanol; acetyl-coenzyme A; niacin; riboflavin; thiamin; ribose; electron transporters such as. . .

ACCESSION NUMBER: 2002:224254 USPATFULL
TITLE: Sunscreen compositions containing a dibenzoylmethane derivative
INVENTOR(S): Cole, Curtis, Ringoes, NJ, United States
Natter, Florence, Hillsborough, NJ, United States
PATENT ASSIGNEE(S): Johnson & Johnson Consumer Companies, Inc., Skillman, NJ, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6444195	B1	20020903
APPLICATION INFO.:	US 2001-883416		20010618 (9)
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	GRANTED		
PRIMARY EXAMINER:	Dodson, Shelley A.		
LEGAL REPRESENTATIVE:	Harriman, Erin M.		

NUMBER OF CLAIMS: 21
EXEMPLARY CLAIM: 1
NUMBER OF DRAWINGS: 0 Drawing Figure(s); 0 Drawing Page(s)
LINE COUNT: 485
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L24 ANSWER 15 OF 17 USPATFULL on STN

SUMM . . . benzoyl peroxide, sulfur resorcinol, ascorbic acid, D-panthenol, hydroquinone, sunscreen agents, anti-inflammatory agents, skin lightening agents, antimicrobial and antifungal agents, estrogens, 2-dimethylaminoethanol, lipoic acid, amino acids such a proline and tyrosine, lactobionic acid, acetyl-coenzyme A, niacin, riboflavin, thiamin, ribose, electron transporters such as NADH and FADH2, botanical extracts such as aloe.

ACCESSION NUMBER: 2002:201667 USPATFULL
TITLE: Cosmetic compositions containing creatine, carnitine, and/or pyruvic acid
INVENTOR(S): Shapiro, Stanley S., Livingston, NJ, United States
Martin, Katharine M., Ringoes, NJ, United States
Shaya, Steven A., Highlands, NJ, United States
Kaminski, Claudia K., Milford, NJ, United States
PATENT ASSIGNEE(S): Johnson & Johnson Consumer Companies, Inc., Skillman, NJ, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6432424	B1	20020813
APPLICATION INFO.:	US 2000-606491		20000629 (9)
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	GRANTED		
PRIMARY EXAMINER:	Moezie, Minna		
ASSISTANT EXAMINER:	Berman, Alycia		
LEGAL REPRESENTATIVE:	McGowen, William E.		
NUMBER OF CLAIMS:	24		
EXEMPLARY CLAIM:	1		
NUMBER OF DRAWINGS:	0 Drawing Figure(s); 0 Drawing Page(s)		
LINE COUNT:	691		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L24 ANSWER 16 OF 17 USPATFULL on STN

SUMM . . . sulfur resorcinol, ascorbic acid, D-panthenol, hydroquinone, sunscreen agents, keratolytic agents, anti-inflammatory agents, skin lightening agents, antimicrobial and antifungal agents, estrogens, 2-dimethylaminoethanol, lipoic acid, amino acids such a proline and tyrosine, lactobionic acid, acetyl-coenzyme A, niacin, riboflavin, thiamin, ribose, electron transporters such as NADH and FADH2, botanical extracts such as aloe.

ACCESSION NUMBER: 2002:191251 USPATFULL
TITLE: Astringent composition and method of use
INVENTOR(S): Watson, Geraldine A., Redondo Beach, CA, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002102314	A1	20020801
APPLICATION INFO.:	US 6482446	B2	20021119
DOCUMENT TYPE:	US 2000-728012	A1	20001201 (9)
FILE SEGMENT:	Utility		
LEGAL REPRESENTATIVE:	Philip S. Johnson, One Johnson & Johnson Plaza, New Brunswick, NJ, 08933-7003		
NUMBER OF CLAIMS:	21		

EXEMPLARY CLAIM:

1

LINE COUNT:

345

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L24 ANSWER 17 OF 17 USPATFULL on STN

DETD . . . benzoyl peroxide, sulfur resorcinol, ascorbic acid, D-panthenol, hydroquinone, sunscreen agents, anti-inflammatory agents, skin lightening agents, antimicrobial and antifungal agents, estrogens, 2-dimethylaminoethanol, lipoic acid, amino acids such as proline and tyrosine, lactobionic acid, acetyl-coenzyme A, niacin, riboflavin, thiamin, ribose, electron transporters such as NADH and FADH₂, botanical extracts such as aloe.

ACCESSION NUMBER: 2002:81526 USPATFULL

TITLE: Method of promoting skin cell metabolism

INVENTOR(S): Shapiro, Stanley S., Livingston, NJ, United States

Martin, Katharine M., Ringoes, NJ, United States

PATENT ASSIGNEE(S): Johnson & Johnson Consumer Companies, Inc., Skillman, NJ, United States (U.S. corporation)

NUMBER	KIND	DATE
--------	------	------

----- ----- -----

PATENT INFORMATION: US 6372791 B1 20020416

APPLICATION INFO.: US 2000-606556 20000629 (9)

DOCUMENT TYPE: Utility

FILE SEGMENT: GRANTED

PRIMARY EXAMINER: Dees, Jose G.

ASSISTANT EXAMINER: George, Konata

LEGAL REPRESENTATIVE: McGowan, William E.

NUMBER OF CLAIMS: 28

EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 0 Drawing Figure(s); 0 Drawing Page(s)

LINE COUNT: 629

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

(FILE 'HOME' ENTERED AT 05:35:24 ON 21 MAR 2005)

FILE 'USPATFULL' ENTERED AT 05:35:29 ON 21 MAR 2005
ACTIVATE L10768359/L

L1 (10638) SEA FILE=USPATFULL ABB=ON PLU=ON TYROSINE AND (LIPOIC OR GLUT
L2 (360) SEA FILE=USPATFULL ABB=ON PLU=ON TYROSINE (100A) (LIPOIC OR G
L3 (174) SEA FILE=USPATFULL ABB=ON PLU=ON L2 AND (TOPICAL OR EXTERNAL)
L4 (198) SEA FILE=USPATFULL ABB=ON PLU=ON TYROSINE (50A) (LIPOIC OR GL
L5 (97) SEA FILE=USPATFULL ABB=ON PLU=ON L4 AND L3
L6 (89) SEA FILE=USPATFULL ABB=ON PLU=ON L5 NOT PERRICONE
L7 (97) SEA FILE=USPATFULL ABB=ON PLU=ON L5 NOT PERRICONE/AU
L8 (10) SEA FILE=USPATFULL ABB=ON PLU=ON (TYROSINE/CLM (50A) (LIPOIC/
L9 (83) SEA FILE=USPATFULL ABB=ON PLU=ON L6 NOT L8

L10 97 S L5
E PERIICONE CICHOMAS/AU
E PERIICONE NICHOLAS/AU
E PERRICONE NICHOLAS/AU
L11 45 S E27-29
L12 90 S L10 NOT L11
L13 90 FOCUS L12 1-
L14 4255 S DIMETHYLAMINOETHANOL
L15 174 S L3
L16 32 S DIMETHYLAMINOETHANOL/AB
L17 269 S DIMETHYLAMINOETHANOL/CLM
L18 273 S L16 OR L17
L19 8 S L18 AND L15
L20 4255 S DIMETHYLAMINOETHANOL
L21 50125 S TYROSINE
L22 26269 S (LIPOIC ACID OR GLUTATHIONE)
L23 23 S L20 (30A) L21 (30A) L22
L24 17 S L23 NOT L11
L25 8 S L20/CLM AND L21/CLM AND L22/CLM
L26 1 S L25 NOT L11
L27 7485 S L20/CLM OR L21/CLM OR L22/CLM
L28 2220 S L20/AB OR L21/AB OR L22/AB
L29 6 S L28 AND L27 AND L23
L30 0 S L29 NOT L11
L31 9 S L28 AND L27 AND L15
L32 2 S L31 NOT L11

=> save all
ENTER NAME OR (END) :end

=> save all temp
ENTER NAME OR (END) :l10768359/l
'L10768359/l' IN USE
A single name cannot be used for two saved items at the same time.
Enter "Y" if you wish to replace the current saved name with a new
definition. Enter "N" if the current saved definition must be
preserved. You may then reenter the SAVE command with a different
saved name. Enter "DISPLAY SAVED" at an arrow prompt (=>) to see a
list of your currently defined saved names.
REPLACE OLD DEFINITION? Y/(N):y
L# LIST L1-L32 HAS BEEN SAVED AS 'L10768359/l'

=>

L12 ANSWER 87 OF 90 USPATFULL on STN

AB A preparation for **external** application to the skin which comprises disodium adenosine triphosphate and tranexamic acid for prevention of skin roughening and skin improvement. . . .

SUMM This invention relates to preparations for **external** application to the skin, more particularly **external** preparations having powerful effects of preventing skin roughening and improving the skin. The **external** preparation of the present invention is suitably applied to cosmetics, such as clear lotions, creams, milky lotions, facial packs, and. . . .

SUMM One of the major purposes of **external** preparations for the skin such as cosmetics consists in prevention of skin roughening and skin improvement. For this purpose, humectants,

SUMM and cosmetics (see JP-B-47-1479, the term "JP-B" as used herein means an "examined published Japanese patent application"). However, preparations for **external** application containing a large amount of tranexamic acid are sticky and feel unpleasant when applied to the skin. Further, ginseng. . . .

SUMM been completed by taking these circumstances into consideration. An object of the present invention is to provide a preparation for **external** application to the skin which produces improved effects on the skin in healing of wounds, prevention of skin roughening, and. . . .

SUMM The present invention relates to a preparation for **external** application to the skin which contains disodium adenosine triphosphate and tranexamic acid.

DRWD acid, sorbic acid, alkyl p-hydroxybenzoates (e.g., ethyl p-hydroxybenzoate or butyl p-hydroxybenzoate), and hexachlorophene; amino acids, e.g., glycine, alanine, valine, leucine, **serine**, threonine, phenylalanine, **tyrosine**, aspartic acid, asparagine, glutamine, taurine, arginine, and histidine, and alkali metal salts and a hydrochloride of these amines; organic acids, e.g., acylsarcosine (e.g., sodium lauroylmethylsarcosine), **glutathione**, malic acid and tartaric acid; vitamins such as vitamin A and its derivatives, vitamin B group and its derivatives including. . . .

DETD Preparations for **external** application to the skin were prepared according to the formulation shown in Tables 1 and 2 and tested for an. . . .

DETD In the following Examples 6 to 13 preparations for **external** application were prepared. All of the preparations exhibited effects of preventing skin roughness and improving the skin conditions without causing. . . .

CLM What is claimed is:

1. A preparation for **external** application to the skin which comprises 0.0005 to 3.0% by weight disodium adenosine triphosphate, 0.01 to 3.0% by weight tranexamic. . . .
2. The preparation for **external** application to the skin as claimed in claim 1, which is for amelioration of skin roughening.

ACCESSION NUMBER: 97:73295 USPATFULL
TITLE: Cosmetic composition
INVENTOR(S): Ogawa, Haruo, Kanagawa, Japan
 Nishiyama, Shoji, Kanagawa, Japan
 Ito, Kenzo, Kanagawa, Japan
PATENT ASSIGNEE(S): Shiseido Company, Ltd., Tokyo, Japan (non-U.S.
corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 5658578		19970819
APPLICATION INFO.:	US 1995-505666		19950721 (8)

	NUMBER	DATE
PRIORITY INFORMATION:	JP 1995-158448	19950601

L19 ANSWER 8 OF 8 USPATFULL on STN

ACCESSION NUMBER: 2001:208908 USPATFULL

TITLE: Topical scar treatments using alkanolamines

INVENTOR(S): Perricone, Nicholas V., 27 Coginchaug Ct., Guilford, CT, United States 06437

NUMBER	KIND	DATE
--------	------	------

PATENT INFORMATION: US 6319942 B1 20011120
APPLICATION INFO.: US 2001-875317 20010606 (9)

DOCUMENT TYPE: Utility

FILE SEGMENT: GRANTED

PRIMARY EXAMINER: Henley, III, Raymond

LEGAL REPRESENTATIVE: Krinsky, Mary M.

NUMBER OF CLAIMS: 20

EXEMPLARY CLAIM: 1

LINE COUNT: 540

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

TI Topical scar treatments using alkanolamines

AB Cutaneous scars are reduced by the topical application of compositions containing an alkanolamine such as ethylaminoethanol, methylaminoethanol, dimethylaminoethanol, isopropanolamine, triethanolamine, isopropanoldimethylamine, ethylethanolamine, 2-butanolamine, choline, serine, and mixtures thereof. Compositions may be applied directly to scar tissue, or embedded in linaments held against the scars. Dimethylaminoethanol in amounts ranging from about 0.1% to about 10% by weight of the total composition is particularly preferred. Adjunct ingredients such as lipoic acid, tyrosine, ascorbyl palmitate, and glycolic acid may be added to scar-reducing formulations, and are desirable in many embodiments.

SUMM . . . Other treatments include application of silicone pads to the scar tissue surface, sometimes under pressure provided by an elastomeric bandage, topical application of silicone gel sheets, with or without added vitamin E (Palmieri, B., et al., J. Derm., 1995, 34: 506-509), and topical or intralesional treatment with corticosteroids.

SUMM It is another and more specific objective of the invention to provide topical compositions and simple methods for scar reduction and inhibition based upon direct topical application of compositions containing active ingredients and/or linaments such as a silicone gel sheet embedded with active ingredients, to scars. . . .

SUMM . . . accomplished by the present invention, which provides compositions and methods for the treatment and/or inhibition of cutaneous scars, which comprises topical application to the scars or injured skin areas of an effective amount of an alkanolamine such as ethylaminoethanol, methylaminoethanol, dimethylaminoethanol, isopropanolamine, triethanolamine, isopropanoldimethylamine, ethylethanolamine, 2-butanolamine, choline, serine, and mixtures thereof. Dimethylaminoethanol is particularly preferred. Amounts of active alkanolamine ingredient in scar-reducing topical compositions of the invention range from about 0.1% to about 10%, more narrowly from about 1% to about 3%, by weight of the total composition. Adjunct ingredients such as lipoic acid, tyrosine, a fatty acid ester of ascorbic acid, e.g., ascorbyl palmitate, and/or an α -hydroxy acid, e.g., glycolic acid may be added to scar-reducing formulations of the invention. One particularly efficacious embodiment for scars employs a composition containing diethylaminoethanol, lipoic acid, and tyrosine; the composition may, optionally, contain other ingredients. Methods and compositions of the invention are particularly efficacious for acne

- scars and. . .
- DETD Methods of the invention involve the **topical** administration of **dimethylaminoethanol** and/or other structurally related alkanolamines, or their biologically equivalent derivatives, to mammalian skin scars for the reduction and inhibition of. . . types of skin trauma. Active alkanolamine active ingredients may be applied alone, or in combination with other ingredients such as **lipoic** acid and/or **tyrosine** to enhance the efficacy of the scar treatment.
- DETD However, only effective amounts of alkanolamines are needed to reduce scars, so generally **topical** application is accomplished in association with a carrier, and particularly one in which the alkanolamine active ingredient is soluble per. . . dermatologically acceptable carrier or vehicle (e.g., as a lotion, cream, ointment, soap, stick, or the like) so as to facilitate **topical** application and, in some cases, provide additional therapeutic effects as might be brought about, e.g., by moisturizing of the affected. . . simple solvent or dispersant such as water, it is generally preferred that the carrier comprise a composition more conducive to **topical** application, and particularly one which will form a film or layer on the skin to which it is applied so. . .
- DETD Whether they are **topical** compositions directly applied to scar tissue or linaments embedded with alkanolamine active ingredients, some embodiments of this invention contain at. . .
- DETD Scar-reducing **topical** compositions of the invention can comprise additional ingredients commonly found in skin care compositions, such as, for example, emollients, skin. . .
- DETD Typical compositions of the invention comprise diethylaminoethanol alone; diethylaminoethanol and **lipoic** acid; a combination of diethylaminoethanol, **lipoic** acid, and **tyrosine**; and a combination of diethylaminoethanol, **lipoic** acid, **tyrosine**, and glycolic acid. Embodiments employing the occlusive effects of silicone pads or gel sheets to diminish scars generally employ higher. . . provide maximal efficacy. A preferred embodiment used in a double blind, placebo-controlled study was a composition containing 3% by weight **dimethylaminoethanol**, 5% **tyrosine**, 3% **lipoic** acid, and 7% glycolic acid.
- CLM What is claimed is:
2. A method according to claim 1 wherein the alkanolamine is selected from the group consisting of ethylaminoethanol, methylaminoethanol, **dimethylaminoethanol**, isopropanolamine, triethanolamine, isopropanoldimethylamine, ethylethanolamine, 2-butanolamine, choline, serine, and mixtures thereof.
 3. A method according to claim 2 wherein the alkanolamine is **dimethylaminoethanol**.
 14. A method for the treatment or inhibition of cutaneous scar tissue comprising applying to said tissue a composition containing from about 0.1% to about 10% by weight of an alkanolamine selected from the group consisting of ethylaminoethanol, methylaminoethanol, **dimethylaminoethanol**, isopropanolamine, triethanolamine, isopropanoldimethylamine, ethylethanolamine, 2-butanolamine, choline, serine, and mixtures thereof.
 15. A method according to claim 14 wherein the composition comprises **dimethylaminoethanol**.
 19. A method for reducing cutaneous scar tissue comprising applying to said tissue a linament embedded with an effective amount of **dimethylaminoethanol**.
- . . . method according to claim 19 wherein the linament is embedded with a

composition containing from about 0.1% to about 10% **dimethylaminoethanol** and at least one other ingredient selected from the group consisting of from about 0.1% to about 7% by weight **lipoic acid**, from about 0.1% to about 5% by weight **tyrosine**, from about 1% to about 10% by weight of glycolic acid, from about 0.5% to about 15% by weight ascorbyl. . .

=>

L19 ANSWER 7 OF 8 USPATFULL on STN
ACCESSION NUMBER: 2002:70014 USPATFULL
TITLE: Treatment of acne using lipoic acid
INVENTOR(S): Perricone, Nicholas V., 27 Coginchaug Ct., Guilford,
CT, United States 06437

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6365623	B1	20020402
APPLICATION INFO.:	US 1999-475514		19991230 (9)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. US 1999-415792, filed on 8 Oct 1999 Continuation-in-part of Ser. No. US 1997-971820, filed on 17 Nov 1997, now patented, Pat. No. US 5965618, issued on 12 Oct 1999		
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	GRANTED		
PRIMARY EXAMINER:	Travers, Russell		
LEGAL REPRESENTATIVE:	Krinsky, Mary M.		
NUMBER OF CLAIMS:	12		
EXEMPLARY CLAIM:	1		
NUMBER OF DRAWINGS:	0 Drawing Figure(s); 0 Drawing Page(s)		
LINE COUNT:	714		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Active acne and acneiform scars are treated by **topical** application of a composition containing lipoic acid and/or a lipoic acid derivative such as dihydrolipoic acid, a lipoic or dihydrolipoic amide, a lipoic or dihydrolipoic acid salt, and mixtures of any of these to reduce erythema, pore size, and scarring. **Topical** application of lipoic acid and/or a lipoic acid derivatives are advantageously used with at least one adjunct ingredient such as . . .
SUMM . . . abnormal keratinization and impaction in the pilosebaceous canal causing obstruction to sebum flow; and (3) proliferation of P. acnes. Thus, **topical** agents that remove comedones, such as **topical** retinoids are particularly effective because they normalize desquamation within the follicular orifice, which allows the sebum to flow freely onto. . . pruritis, burning/stinging, and scaling/flaking (Physicians' Desk Reference®, 54th ed. 2000, pp 502-503, 1104-1105, and 2139-2142, hereinafter referred to as "PDR"). **Topical** vitamin A preparations and benzoyl peroxide have been used to treat acne for some time. However, it has been recently. . . thickness, and deleterious changes in elastin and glycosaminoglycan content (Ibbotson, S. H., et al., J. Inves. Derm., 1999, 112: 933-938). **Topical** and oral antibiotics (especially tetracycline, erythromycin, and clindamycin) are sometimes prescribed for patients with inflammatory papules and pustules; but, in. . .
SUMM . . . been traditionally treated with invasive methods such as scar revision, laser ablation, and chemical peels. Non-invasive techniques have consisted of **topical** application of tretinoin, as well as the application of estrogens and α -hydroxy acids. None of these non-invasive procedures have been. . .
SUMM . . . patent publication (JP 63008315), lipoic acid in cosmetics at concentrations of 0.01% to 1%, preferably 0.05% to 0.5%, or in **topical** "quasi-drugs" at concentrations of 0.1% to 1.5%, preferably 0.5% to 1.0%, were suggested for inhibiting tyrosinase, and thus melanin formation,. . .
SUMM . . . intravenous, or infusions (column 3, lines 28 to 30, 51, 62 to 63 and 65), but solutions and emulsions for **topical** application were mentioned (column 6, lines 29 to 34 and 65 to 68, and column 8, lines 16 to 18).. . .
SUMM It is another and more specific objective of the invention to provide **topical** compositions and methods for acne lesion and acne scar treatment based upon the application of compositions containing lipoic

SUMM acid and/or . . . invention, which provides compositions and methods for the treatment of acne vulgaris, and improvements of currently employed therapies, which comprise **topical** application to skin areas exhibiting acne of an effective amount of **lipoic** acid, **lipoic** acid derivatives or mixtures thereof, typically in association with a dermatologically acceptable carrier. In most preferred embodiments, at least one. . . to, α -hydroxy acids such as glycolic and/or lactic acid; tocotrienols; fatty acid esters of ascorbic acid such as ascorbyl palmitate; **tyrosine**; antibiotics such as erythromycin, clindamycin, or tetracycline; retinoids such as tretinoin, adapalene, or tazarotene; or methyl- or ethyl-aminoalcohols such as **dimethylaminoethanol**. Benzoyl peroxide is included in some compositions. Adjunct ingredients enhance the efficacy of the treatment, and minimize or eliminate skin. . .

SUMM However, only effective amounts of lipoic acid are needed to treat acne and acneiform scars, so generally **topical** application to exposed or affected skin sites is accomplished in association with a carrier, and particularly one in which the. . .

SUMM . . . of a relatively simple solvent or dispersant, it is generally preferred that the carrier comprise a composition more conducive to **topical** application, and particularly one which will form a film or layer on the skin to which it is applied so. . .

SUMM . . . their effects, but minimizes or eliminates their side effects. Adjunct ingredients include, but are not limited to, not only retinoids, **topical** antibiotics, and benzoyl peroxide conventionally used in acne treatments, but also methyl-/ethyl-aminoalcohols, α -hydroxy acids, **tyrosine** tocotrienols, and fatty acid esters. . .

SUMM . . . provides a method for treating acne using less retinoid than would be required if a retinoid is used alone, because **topical** application of retinoids results in skin irritation in some patients. As set out in the PDR sections cited above, even. . .

SUMM Lipoic acid may also be used in combination with **topical** or oral antibiotics such as tetracycline, clindamycin, and erythromycin sometimes used for acne cases, particularly for patients with inflammatory papules. . .

SUMM It is an advantage of the invention that **topical** application of **lipoic** acid provides a simple, non-invasive, nontoxic, over-the-counter **topical** method for treating all phases of acne. **Lipoic** acid compositions decrease erythema observed with acne pustules, papules and whiteheads, and a marked decrease in lesion numbers. The effect is enhanced by use of adjunct ingredients such as **dimethylaminoethanol**, α -hydroxy acids, and/or **tyrosine**. **Lipoic** acid compositions decrease pore size, minimizing sebum accumulation and keratinous debris that cause both whiteheads and blackheads observed in acne. **Lipoic** acid minimizes scar formation, and provides marked losses of scar borders and decreases in scar depth where scars have already formed. Topically applied **lipoic** acid also seems to fill in scar tissue, making it more equal to adjacent normal skin. Moreover, compositions containing **lipoic** acid with adjunct ingredients such as retinoids, α -hydroxy acids, **tyrosine**, and/or **dimethylaminoethanol**, appear to successfully treat active acne lesions without harming surrounding skin tissue. And with these physical effects, persons using **lipoic** acid **topical** compositions experience a reduction in the social and psychological stress often associated with acne patients suffering facial disfigurements.

CLM What is claimed is:

according to claim 1 wherein the composition further comprises a methyl- or ethyl-aminoalcohol ingredient selected from the group consisting of **dimethylaminoethanol**, monomethylaminoethanol, diethylaminoethanol, monoethylaminoethanol, their propanol and butanol

counterparts, derivatives acylated with organic acids, and mixtures thereof.

12. A method according to claim 11 wherein the aminoalcohol ingredient is **dimethylaminoethanol**.

L19 ANSWER 6 OF 8 USPATFULL on STN

ACCESSION NUMBER: 2002:224254 USPATFULL
TITLE: Sunscreen compositions containing a dibenzoylmethane derivative
INVENTOR(S): Cole, Curtis, Ringoes, NJ, United States
Natter, Florence, Hillsborough, NJ, United States
PATENT ASSIGNEE(S): Johnson & Johnson Consumer Companies, Inc., Skillman, NJ, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6444195	B1	20020903
APPLICATION INFO.:	US 2001-883416		20010618 (9)
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	GRANTED		
PRIMARY EXAMINER:	Dodson, Shelley A.		
LEGAL REPRESENTATIVE:	Harriman, Erin M.		
NUMBER OF CLAIMS:	21		
EXEMPLARY CLAIM:	1		
NUMBER OF DRAWINGS:	0 Drawing Figure(s); 0 Drawing Page(s)		
LINE COUNT:	485		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

DETD . . . tocopheryl acetate; retinoids such retinol, retinal, retinyl palmitate, retinyl acetate, and retinoic acid; hormones such as estrogens and dihydroxyandrostene dione; **2-dimethylaminoethanol**; **lipoic** acid; amino acids such a proline and **tyrosine**; lactobionic acid; self-tanning agents such as dihydroxy acetone; dimethyl aminoethanol; acetyl-coenzyme A; niacin; riboflavin; thiamin; ribose; electron transporters such as. . .
DETD . . . antioxidants, preservatives, and chelating agents are listed in pp. 1612-13, 1626, and 1654-55 of the ICI Handbook. In addition, the **topical** compositions useful herein can contain conventional cosmetic adjuvants, such as dyes, opacifiers (e.g., titanium dioxide), pigments, and fragrances.
DETD . . . the skin or hair of a human. The cosmetic compositions useful in the subject invention, thus, involve formulations suitable for **topical** application to mammalian skin or hair, the formulation comprising (i) dibenzoylmethane derivative(s), (ii) a diester or polyester of a naphthalene. . . compounds/agents such as the other UV-A or UV-B absorbers/reflectors listed herein, and/or other cosmetically active agents and (v) a cosmetically-acceptable **topical carrier**. The term "cosmetically-acceptable **topical carrier**" refers to a carrier for **topical** use that is capable of having the dibenzoylmethane, the diester or polyester of a naphthalene dicarboxylic acid, the benzophenone derivative. . .
DETD The **topical** compositions useful in the present invention may be used for a variety of cosmetic uses, including, but not limited to,. . .
CLM What is claimed is:
. . . hydroxy acids, benzoyl peroxide, sulfur resorcinol, D-panthenol, hydroquinone, anti-inflammatory agents, skin lightening agents, antimicrobial agents, antifungal agents, vitamins, retinoids, hormones, **2-dimethylaminoethanol**, lipoic acid, amino acids, lactobionic